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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|---|---------------|----------------------|---------------------|------------------|
| 10/517,171 | 06/28/2005 | Bernd Schumann | 10191/3924 | 7411 |
| 26646 | 7590 | 11/25/2008 | EXAMINER | |
| KENYON & KENYON LLP ONE BROADWAY NEW YORK, NY 10004 | | | SALZMAN, KOURNEY R | |
| ART UNIT | PAPER NUMBER | 1795 | | |
| MAIL DATE | DELIVERY MODE | 11/25/2008 | PAPER | |

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

| | | | |
|------------------------------|------------------------|---------------------|--|
| Office Action Summary | Application No. | Applicant(s) | |
| | 10/517,171 | SCHUMANN, BERND | |
| | Examiner | Art Unit | |
| | KOURTNEY R. SALZMAN | 1795 | |

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

1) Responsive to communication(s) filed on 15 August 2008.
 2a) This action is **FINAL**. 2b) This action is non-final.
 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

4) Claim(s) 16-31 is/are pending in the application.
 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
 5) Claim(s) _____ is/are allowed.
 6) Claim(s) 16-31 is/are rejected.
 7) Claim(s) _____ is/are objected to.
 8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

9) The specification is objected to by the Examiner.
 10) The drawing(s) filed on _____ is/are: a) accepted or b) objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 a) All b) Some * c) None of:
 1. Certified copies of the priority documents have been received.
 2. Certified copies of the priority documents have been received in Application No. _____.
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

1) Notice of References Cited (PTO-892)
 2) Notice of Draftsperson's Patent Drawing Review (PTO-948)
 3) Information Disclosure Statement(s) (PTO/SB/08)
 Paper No(s)/Mail Date _____.
 4) Interview Summary (PTO-413)
 Paper No(s)/Mail Date _____.
 5) Notice of Informal Patent Application
 6) Other: _____.

DETAILED ACTION

Summary

1. The arguments filed August 15, 2008 have been fully considered.
2. Claims 16-31 are currently pending.
3. The USC 103 rejections of claims 16-31 from the previous office action are withdrawn.

Claim Rejections - 35 USC § 103

4. The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.
5. Claims 16-27 and 29-31 are rejected under 35 U.S.C. 103(a) as being unpatentable over YAMADA et al (US 6,210,641) and KUNIMOTO et al (US 2002/0017461), in view of KATO et al (US 5,492,190).

YAMADA et al teaches a gas sensor used in a combustion engine. In figure 53B YAMADA et al teaches a cavity A within the electrolytic pieces 91-93. The inner electrode of the pumping cell is found to be 91b, while the outer electrode is 91a found outside the cavity and electrolyte. There are two inlets or diffusion channels, which allow the exhaust gas to feed to chamber A. To reach the cavity, the exhaust gas flows through a catalyst layer, 17, not contained in a chamber.

YAMADA et al fails to teach an explicit prechamber containing a catalyst (functioning as a catalytic converter) and diffusion channel which feeds from the prechamber to the cavity.

KUNIMOTO et al discloses a similar gas sensor containing a cavity 3 with a pumping cell 33 in figure 1. KUNIMOTO et al teaches a prechamber 5, fed exhaust gas through 20. In paragraph 38 KUNIMOTO et al teaches a diffusion hole 6 can be used to control the inflow of gas to the prechamber from the cavity 3. The porous member 25 in the prechamber functions as a catalytic converter for reducing the incoming exhaust gas.

At the time of invention, it would have been obvious to one of ordinary skill in the art to utilize the prechamber packing of KUNIMOTO et al in the oxygen sensor of YAMADA et al because the organization of the catalytic treatment in a prechamber allows for accurate readings as the gases are more completely reduced, where simply flowing through the catalyst has led to insufficient reduction of gases and therefore interfering gas species during measurement. (KUNIMOTO et al paragraphs 7-9)

The combination of YAMADA et al and KUNIMOTO et al fails to teach the diffusion barrier to be present within the diffusion channel.

KATO et al teaches a gas sensor utilizing a porous member 58, or diffusion barrier, to control gas diffusion from one cavity to another within a channel as defined by the electrolytes 52f and 52d.

At the time of the invention, it would have been obvious to one of ordinary skill to use the diffusion barrier of KATO et al in substitution for the diffusion hole of KUNIMOTO et al in the sensor of YAMADA et al, or to place the diffusion barrier of KATO et al within the diffusion hole of KUNIMOTO et al in the sensor of YAMADA et al because KATO et al teaches in column 9, lines 20-24, a diffusion barrier passage can be a porous member, as shown in the KATO et al reference number 58, or the diffusion hole as taught by KUNIMOTO et al reference number 6. Therefore, the combination of the porous barrier material and the physical barrier of the diffusion hole would also be an obvious combination.

Regarding claim 17, YAMADA et al teaches the use of an alumina catalyst (c. 14, l. 21-23) which will oxidize the incoming hydrogen gas. (c. 13, l. 59-61)

Regarding claim 18, figure 1 of KUNIMOTO et al shows the catalyst to fill the entirety of the chamber.

Regarding claims 19 and 20, YAMADA et al also shows the use of inside and outer platinum electrodes covered in catalyst indicating the use of the electrodes for electrochemical catalysis. (c. 11, l. 11-43) KUNIMOTO et al also discloses use of numerous precious metal electrodes in paragraph 64.

Regarding claim 21, these electrodes are explicitly used for oxidation and are shown to be present in figure 3 of KUNIMOTO et al across the chamber as reference number 17.

Regarding claim 22, figure 3 of KUNIMOTO et al shows all the entrances and chamber walls which run the length of the sensor to be substantially parallel.

Regarding claim 23, the center axes of the diffusion channel and the intake opening are shown to be parallel in the figure 1 of KUNIMOTO et al.

Regarding claim 24, figure 1 of KUNIMOTO et al also shows the diffusion passageway 6 to be of a much smaller relative cross sectional area than the prechamber 5.

Regarding claim 25, KUNIMOTO et al teaches the use of direct current power supply as reference numbers 33 and 34 and show in figure 3 to be permanently connected. Circuit 34 moves current from the site of the 34 to the inner electrode to the outer electrode. Due to all the resistance the current will encounter as it moves through the circuit, the voltage would be higher at the source, or at the input, than after flowing through the electrodes.

Regarding claim 26, YAMADA et al teaches a porous ceramic like alumina and a catalytic metal, like Pt-Rh or Pt. A cermet is known in the art to be a mix of a ceramic and metal. Here, the coating is shown to be around the electrodes. (column 19, lines 35-52)

Regarding claim 27, weight is given to the preamble when it discloses structural limitations therefore, in figure 53B YAMADA et al teaches a cavity A within the electrolytic pieces 91-93. The inner electrode of the pumping cell is found to be 91b, while the outer electrode is 91a found outside the cavity and electrolyte. There are two inlets or diffusion channels, which allow the exhaust gas to feed to chamber A. To reach the cavity, the exhaust gas flows through a catalyst layer, 17, not contained in a chamber.

YAMADA et al fails to teach an explicit prechamber containing a catalyst (functioning as a catalytic converter) and diffusion channel which feeds from the prechamber to the cavity.

KUNIMOTO et al discloses a similar gas sensor containing a cavity 3 with a pumping cell 33 in figure 1. KUNIMOTO et al teaches a prechamber 5, fed exhaust gas through 20. In paragraph 38, KUNIMOTO et al discloses the gas diffusion hole 6 can be used to control the inflow of gas to the prechamber from

the cavity 3. The porous member 25 in the prechamber functions as a catalytic converter for reducing the incoming exhaust gas.

At the time of invention, it would have been obvious to one of ordinary skill in the art to utilize the prechamber packing of KUNIMOTO et al in the oxygen sensor of YAMADA et al because the organization of the catalytic treatment in a prechamber allows for accurate readings as the gases are more completely reduced, where simply flowing through the catalyst has led to insufficient reduction of gases and therefore interfering gas species during measurement. (KUNIMOTO et al paragraphs 7-9)

The combination of YAMADA et al and KUNIMOTO et al fails to teach the diffusion barrier to be present within the diffusion channel.

KATO et al teaches a gas sensor utilizing a porous member 58, or diffusion barrier, to control gas diffusion from one cavity to another within a channel as defined by the electrolytes 52f and 52d.

At the time of the invention, it would have been obvious to one of ordinary skill to use the diffusion barrier of KATO et al in substitution for the diffusion hole of KUNIMOTO et al in the sensor of YAMADA et al, or to place the diffusion barrier of KATO et al within the diffusion hole of KUNIMOTO et al in the sensor of

YAMADA et al because KATO et al teaches in column 9, lines 20-24, a diffusion barrier passage can be a porous member, as shown in the KATO et al reference number 58, or the diffusion hole as taught by KUNIMOTO et al reference number 6. Therefore, the combination of the porous barrier material and the physical barrier of the diffusion hole would also be an obvious combination.

Regarding claim 29, as oxidation of hydrogen is the goal of an oxygen sensor, and atomic hydrogen is known to burn above a temperature of 600°C, as stated in c. 15, lines. 64-66 of YAMADA et al, any operation temperature above this temperature would be obvious to one of ordinary skill in the art.

Regarding claim 30, the response time of the system based on the amount of catalyst used is shown in figure 32 of YAMADA et al. Points of the graph show operation far exceeding one minute.

Regarding claim 31, due to the amount of testing required to maintain the air-fuel ratio control system of the catalytic converter of YAMADA et al it would be obvious to repeat the application of voltage for the duration of the engine's requirement for air-fuel ratio calculation. (c. 11, l. 55-c.12, l.13)

6. Claim 28 is rejected under 35 U.S.C. 103(a) as being unpatentable over YAMADA et al (US 6,210,641 B1), KUNIMOTO et al (US 2002/0017461 A1) and KATO

et al (US 5,942,190) as applied to claim 27 above, and further in view of NADANAMI et al (US 2002/0092780).

Regarding claim 28, the combination of YAMADA et al and KUNIMOTO et al teaches all the limitations of claim 27.

The combination YAMADA et al and KUNIMOTO et al fails to teach the application of a DC voltage higher than that of the decomposition voltage of the solid electrolyte.

NADANAMI et al teaches the application of a voltage at the level of decomposition of the electrolyte in the sensor in paragraph 17.

It would be obvious to one of ordinary skill in the art to raise the voltage above that of the decomposition voltage if voltage at that level is currently present.

At the time of invention, it would obvious to apply the DC voltage of NADANAMI et al to the sensor of YAMADA et al and KUNIMOTO et al because it causes the successful transport of protons through the sensor, limiting proton current flow, a goal of sensors in the industry.

Response to Arguments

7. Applicant's arguments, see the 3rd-5th fully paragraphs of the second section of the remarks, filed August 15, 2008, with respect to the rejection(s) of claim(s) 16 and 27

under 35 USC 103 have been fully considered and are persuasive. Therefore, the rejection has been withdrawn. However, upon further consideration, a new ground(s) of rejection is made in view of a different interpretation of the previous reference KUNIMOTO et al and the newly applied reference KATO et al. KUNIMOTO et al is still used to teach the prechamber containing the catalytic converter. KUNIMOTO et al also still teaches a diffusion channel, reference number 6. KATO et al is used to teach the use of a porous material or diffusion member, within an electrolytic channel, as the diffusion barrier.

Conclusion

8. Any inquiry concerning this communication or earlier communications from the examiner should be directed to KOURTNEY R. SALZMAN whose telephone number is (571)270-5117. The examiner can normally be reached on Monday to Thursday 6:30AM-5PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Nam Nguyen can be reached on (571) 272-1342. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Kaj K Olsen/
Primary Examiner, Art Unit 1795

krs
11/18/2008